## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

- 1-4. (Canceled)
- 5. (Currently amended) A method of testing for a cedar pollen allergy <u>in a</u> human subject, said method comprising the steps of:
  - (a) preparing T cells from the subjecta subject,
  - (b) preparing an RNA sample from said T cells,
- (c) conducting hybridization with said RNA sample using a probe consisting of the complement of a segment of SEQ ID NO:1, optionally wherein the probe is linked to a label or vector nucleic acideomprising nucleotides of gene 513, wherein the nucleotides consist of a segment of nucleotides between positions 36 and 1171 of SEQ ID NO:1 or the complement thereof wherein said probe is labeled, and
- (d) measuring the amount of RNA that is derived from said subject and that hybridizes with said probe, and comparing said amount with the amount of RNA of a control group which has 3.5 AU/mL or less of cedar pollen specific IgE, wherein if the amount of gene 513 RNA is significantly higher in a sample from the subject than in the control group, then the subject is determined to have a cedar pollen allergy.
- 6. (Currently amended) A method of testing for a cedar pollen allergy in a human subject, said method comprising the steps of:
  - (a) preparing T cells from the subject a subject,
  - (b) preparing an RNA sample from said T cells,
- (c) synthesizing cDNA by conducting reverse transcription reaction with said RNA sample,

- (d) conducting polymerase chain reaction (PCR) using said cDNA as template and a primer consisting of a segment of gene 513 SEQ ID NO:1 or the complement thereof, optionally wherein the probe is linked to a label or vector nucleic acid-that comprising nucleotides of gene 513, wherein the nucleotides consist of a segment of nucleotides between positions 36 and 1171 of SEQ ID NO: 1 or the complement thereof, and
- (e) comparing the amount of DNA amplified by said PCR with a control group which has 3.5 AU/mL or less of cedar pollen specific IgE, wherein if the amount of gene 513 cDNA is significantly higher in a sample from the subject than in the control group, then the subject is determined to have a cedar pollen allergy.
- 7. (Original) The method of claim 6, wherein said PCR is carried out by a PCR amplification monitoring method.
- 8. (Currently amended) The method of <u>claim</u> elaims 5, wherein said T cells are prepared from peripheral blood of said subject.
  - 9. (Canceled)

10-20. (Canceled)

- 21. (Currently amended) The method of claim 5, wherein the probe consists of at least 15 nucleotides between positions 36 and 1171 of SEQ ID NO:1 and or the complement thereof.
- 22. (Currently amended) The method of claim 6, wherein the primer consists of at least 15 nucleotides-between positions 36 and 1171 of SEQ ID NO:1 and or the complement thereof.